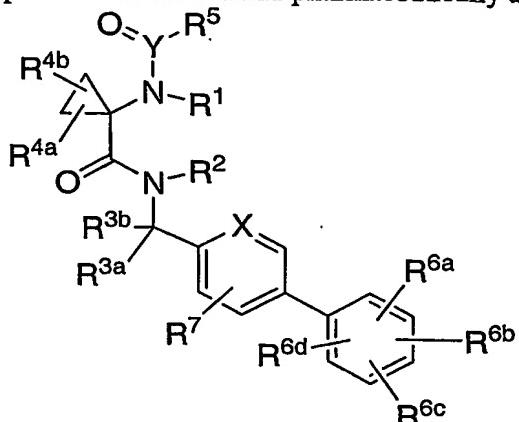


WHAT IS CLAIMED IS:

1. A compound of formula I and pharmaceutically acceptable salts thereof:



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wherein

R¹ and R² are independently selected from hydrogen and C₁-4 alkyl;

R^{3a} and R^{3b} are independently selected from hydrogen and C₁-4 alkyl optionally substituted with 1 to 5 halogen atoms;

- 10 R^{4a} and R^{4b} are independently selected from hydrogen, halogen, and C₁-4 alkyl optionally substituted with 1 to 4 groups selected from halogen, OR^a, OC(O)R^a, S(O)_kR^d, OS(O)₂R^d, and NR¹R², or R^{4a} and R^{4b} together with the carbon atom to which they are both attached form an exo-cyclic methylene optionally substituted with 1 to 2 groups selected from C₁-4 alkyl optionally substituted with 1-5 halogens and C₁-4 alkyloxy;
- 15 R⁵ is selected from (1) C₁-6 alkyl optionally substituted with 1 to 5 groups independently selected from halogen, nitro, cyano, OR^a, SR^a, COR^a, SO₂R^d, CO₂R^a, OC(O)R^a, NR^bRC, NR^bC(O)R^a, NR^bC(O)R^a, C(O)NR^bRC, C₃-8 cycloalkyl, (2) C₃-8 cycloalkyl optionally substituted with 1 to 5 groups independently selected from halogen, nitro, cyano and phenyl, (3) C₃-6 alkynyl, (4) C₂-6 alkenyl optionally substituted with hydroxyethyl, (5) (CH₂)_k-aryl optionally substituted with 1 to 3 groups
- 20 independently selected from halogen, nitro, cyano, OR^a, SR^a, C(O)R^a, C₁-4 alkyl and C₁-3 haloalkyl; (6) (CH₂)_k-heterocycle optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, OR^a, SR^a, C₁-4 alkyl and C₁-3 haloalkyl wherein said heterocycle is selected from (a) a 5-membered heteroaromatic ring having a ring heteroatom selected from N, O and S, and optionally having up to 3 additional ring nitrogen atoms wherein said ring is optionally benzo-fused; (b) a 6-membered heteroaromatic ring containing from 1 to 3 ring nitrogen atoms and N-oxides thereof, wherein said ring is optionally benzo-fused; and (c) a 5- or 6-membered non-aromatic heterocyclic ring selected from

- tetrahydrofuryl, 5-oxotetrahydrofuryl, 2-oxo-2H-pyranyl, 6-oxo-1,6-dihdropyridazinyl, (7) C(O)₂R^a, and (8) C(O)NR^bRC;
- R^{6a} is selected from (1) -OSO₂R⁸, (2) -NR^{8a}SO₂R⁹, and (3) -C(R^{8b})(R^{8c})SO₂R⁹;
- R^{6b}, R^{6c}, and R^{6d} are independently selected from (1) hydrogen, (2) halogen, (3) OSO₂R⁸, (4) C₁₋₄ alkyl optionally substituted with 1 to 5 halogen atoms, (5) cyano, (6) nitro, (7) OR^a, and (8) CO₂R^a, or when attached to adjacent carbon atoms R^{6c} and R^{6d} together with the carbon atoms to which they are attached form a 5- to 8-membered saturated or unsaturated ring;
- R⁷ is selected from (1) hydrogen, (2) halogen, (3) cyano, (4) nitro, (5) OR^a, (6) CO₂R^a, (7) C(O)NR^bRC, and (8) C₁₋₄ alkyl optionally substituted with 1 to 5 halogen atoms,
- 10 R⁸ is selected from (1) C₁₋₄ alkyl optionally substituted with 1 to 5 halogen atoms, (2) (CH₂)_k-aryl optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, NR^aC(O)R^a, OR^a, SR^a, CO₂R^a, C₁₋₄ alkyl, C₁₋₃ haloalkyl and NR^bRC, (3) NR^bRC, and (4) hydrogen;
- R^{8a} is selected from hydrogen, C₁₋₄ alkyl optionally substituted with 1 to 5 halogen atoms, halogen, and CO₂R^a, or
- 15 when R^{6a} and R^{6b} are attached to adjacent atoms, R^{8a} and R^{6b} together complete 5- or 6-membered ring;
- R^{8b} and R^{8c} are independently selected from hydrogen, C₁₋₄ alkyl optionally substituted with 1 to 5 halogen atoms, halogen, cyano, nitro, CO₂R^a, and OR^a;
- R⁹ is selected from (1) C₁₋₄ alkyl optionally substituted with 1 to 5 halogen atoms, (2) aryl optionally
- 20 substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, NR^aC(O)R^a, OR^a, SR^a, CO₂R^a, C₁₋₄ alkyl and C₁₋₃ haloalkyl, and (3) (CH₂)_k-aryl optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, NR^aC(O)R^a, OR^a, SR^a, C(O)₂R^a, C₁₋₄ alkyl and C₁₋₃ haloalkyl, or
- R^{8a} and R⁹ together with the atoms to which they are attached form a 5- to 8-membered heterocyclic ring;
- 25 R^a, R^b and R^c are independently selected from (1) hydrogen, (2) C₁₋₄ alkyl optionally substituted with 1 to 5 halogen atoms, (3) phenyl optionally substituted with 1 to 3 groups selected from halogen, cyano, nitro, OH, C₁₋₄ alkyloxy, C₃₋₆ cycloalkyl and C₁₋₄ alkyl optionally substituted with 1 to 5 halogen atoms, and (4) C₃₋₆ cycloalkyl, or
- 30 R^b and R^c together with the nitrogen atom to which they are attached form a 4-, 5-, or 6-membered ring optionally containing an additional heteroatom selected from N, O, and S; or
- R^b and R^c together with the nitrogen atom to which they are attached form a cyclic imide;
- R^d is selected from (1) C₁₋₄ alkyl optionally substituted with 1 to 5 halogen atoms, (2) C₁₋₄ alkyloxy, (3) phenyl optionally substituted with 1 to 3 groups selected from halogen, cyano, nitro, OH, C₁₋₄

alkyloxy, C₃-6 cycloalkyl and C₁-4 alkyl optionally substituted with 1 to 5 halogen atoms, and (4) hydrogen;
 X is selected from CH and N;
 Y is selected from C and S=O; and
 5 k is selected from 0, 1, and 2.

2. A compound of Claim 1 wherein R⁵ is selected from pyrimidinyl and C₁-6 alkyl optionally substituted with 1 to 5 groups independently selected from halogen.

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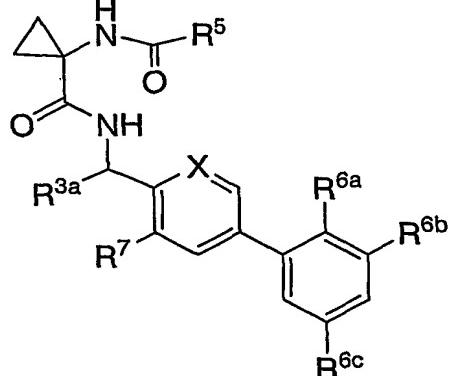
3. A compound of Claim 1 wherein Y is C.

4. A compound of Claim 1 wherein R^{6a} is OSO₂R⁸ and R⁸ is selected from 2,2,2,-trifluoroethyl, trifluoromethyl, methyl, ethyl, propyl, isopropyl, phenyl, benzyl, and dimethylamino; or
 15 R^{6a} is NHSO₂R⁹ and R⁹ is methyl or trifluoromethyl.

5. A compound of Claim 1 wherein R^{6b} is selected from hydrogen, fluorine, and chlorine.

20

6. A compound of Claim 1 having the formula I(2):



I(2)

wherein X is N or CH, R^{3a} is H or C₁-4alkyl, R⁷ is hydrogen or halogen, and R⁵, R^{6a}, R^{6b} and R^{6c} have the same definitions as provided in Claim 1.

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7. A compound of Claim 6 wherein R^{6a} is NHSO₂R⁹; R⁹ is C₁₋₄alkyl optionally substituted with 1 to 5 halogen atoms, R^{6b} is halogen, and R^{6c} is hydrogen or halogen.

8. A compound of Claim 10 wherein R^{6a} is OSO₂R⁸; R⁸ is selected from methyl, 5 trifluoromethyl, ethyl, propyl, isopropyl, benzyl, dimethylamino, 2,2,2-trifluoroethyl, and phenyl; R^{6b} is hydrogen or halogen, and R^{6c} is hydrogen or halogen.

9. A compound of Claim 10 wherein R⁵ is pyrimidinyl or C₁₋₄alkyl optionally substituted with 1 to 5 groups independently selected from halogen.

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10. A compound selected from

3,3'-difluoro-4'-{[(1-[(pyrimidin-5-ylcarbonyl)amino]cyclopropyl)carbonyl]amino]methyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,

3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino]ethyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,

3,3'-difluoro-4'-((1R)-1-{[(1-[(trifluoromethyl)sulfonyl]amino)cyclopropyl]carbonyl]amino}ethyl)-1,1'-biphenyl-2-yl trifluoromethanesulfonate,

1-([(1R)-1-(3,3'-difluoro-2'-[(trifluoromethyl)sulfonyloxy]-1,1'-biphenyl-4-yl)ethyl]amino)carbonyl)-cyclopropanaminium trifluoroacetate,

20 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino]ethyl}-1,1'-biphenyl-2-yl methanesulfonate,

5-chloro-3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,

25 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl ethanesulfonate,

3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl propane-1-sulfonate

3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl propane-2-sulfonate,

30 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl benzenesulfonate,

3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl phenylmethanesulfonate

3,3'-difluoro-4'-(*(1R)*-1-[{({1-[(trifluoroacetyl)amino]cyclopropyl}carbonyl)amino]ethyl}-1,1'-biphenyl-2-yl dimethylsulfamate,

3,3'-difluoro-4'-(*(1R)*-1-[{({1-[(trifluoroacetyl)amino]cyclopropyl}carbonyl)amino]ethyl}-1,1'-biphenyl-2-yl 2,2,2-trifluoroethanesulfonate,

5 3-chloro-3'-fluoro-4'-(*(1R)*-1-[{({1-[(trifluoroacetyl)amino]cyclopropyl}carbonyl)amino]ethyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,

3'-fluoro-4'-(*(1R)*-1-[{({1-[(trifluoroacetyl)amino]cyclopropyl}carbonyl)amino]ethyl}-2-{[(trifluoromethyl)sulfonyl]oxy}-1,1'-biphenyl-3-yl trifluoromethanesulfonate,

10 N-(1-{[({(1R)-1-{3,3'-difluoro-2'-(methyl(methylsulfonyl)amino]-1,1'-biphenyl-4-yl}ethyl)amino]- carbonyl}cyclopropyl)pyrimidine-5-carboxamide,

N-(1-{[({3,3'-difluoro-2'-(methylsulfonyl)amino]-1,1'-biphenyl-4-yl}methyl)amino]carbonyl}- cyclopropyl)pyrimidine-5-carboxamide,

N-{1-[{[2'-(1,1-dioxido-1,2-thiazinan-2-yl)-3,3'-difluoro-1,1'-biphenyl-4-yl]methyl}amino]carbonyl}- cyclopropyl)pyrimidine-5-carboxamide,

15 N-[*(1R)*-1-(3,3'-difluoro-2'-{[(trifluoromethyl)sulfonyl]methyl}-1,1'-biphenyl-4-yl)ethyl]-1-[{(trifluoroacetyl)amino]cyclopropanecarboxamide,

N-[*(1R)*-1-(3,3'-difluoro-2'-{[(trifluoromethyl)sulfonyl]amino}-1,1'-biphenyl-4-yl)ethyl]-1-[{(trifluoroacetyl)amino]cyclopropanecarboxamide, and

20 N-(1-{[({(1R)-1-{3,3'-difluoro-2'-(methylsulfonyl)amino]-1,1'-biphenyl-4-yl}ethyl)amino]carbonyl}- cyclopropyl)pyrimidine-5-carboxamide.

11. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1 and a pharmaceutically acceptable carrier.

25 12. Use of a compound of Claim 1 or a pharmaceutically acceptable salt thereof in the manufacture of a medicament for the treatment or prevention of pain and inflammation.

13. Use of Claim 30 wherein said pain is postherpetic neuropathy, osteoarthritis pain, or dental pain.